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Seini Mantangi

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Patent

Attorney's Docket No. 032405-018

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

UTILITY PATENT
APPLICATION TRANSMITTAL LETTER

Box PATENT APPLICATION

Assistant Commissioner for Patents
 Washington, D.C. 20231

Sir:

Enclosed for filing is the utility patent application of STEPHEN YENCHO and JAIME VARGAS for IMPLANTABLE SUPERELASTIC ANASTOMOSIS DEVICE.

Also enclosed are:

[X] 10 sheet(s) of [X] formal [] informal drawing(s);

[X] An [] executed [X] unexecuted declaration of the inventor(s)

[X] also is enclosed [] will follow.

[X] Small entity status is hereby claimed.



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(10/00)



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09687216-101200

☒ The filing fee has been calculated as follows:

| C L A I M S | | | | | |
|--|------------------|------------|-----------------|-------------------|-----------------|
| | NO. OF CLAIMS | | EXTRA CLAIMS | RATE | FEE |
| Basic Application Fee | | | | | \$710.00 (101) |
| Total Claims | 19 | MINUS 20 = | 0 | × \$18.00 (103) = | .00 |
| Independent Claims | 3 | MINUS 3 = | 0 | × \$80.00 (102) = | .00 |
| If multiple dependent claims are presented, add \$270.00 (104) | | | | | |
| Total Application Fee | | | | | \$710.00 |
| If small entity status is claimed, subtract 50% of Total Application Fee | | | | | (\$355.00) |
| Add Assignment Recording Fee \$ if Assignment document is enclosed | | | | | |
| TOTAL APPLICATION FEE DUE | | | | | \$355.00 |

- ☐ This application is being filed without a filing fee. Issuance of a Notice to File Missing Parts of Application is respectfully requested.
- ☒ A check in the amount of \$ 355.00 is enclosed for the fee due.
- ☐ Charge \$ _____ to Deposit Account No. 02-4800 for the fee due.
- ☒ The Commissioner is hereby authorized to charge any appropriate fees under 37 C.F.R. §§ 1.16, 1.17 and 1.21 that may be required by this paper, and to credit any overpayment, to Deposit Account No. 02-4800. This paper is submitted in duplicate.

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Respectfully submitted,

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Date: October 11, 2000

By: 

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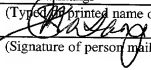
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Seini Mantangi

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PATENT
Attorney Docket No. 032405-018

IMPLANTABLE SUPERELASTIC ANASTOMOSIS DEVICE

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IMPLANTABLE SUPERELASTIC ANASTOMOSIS DEVICE

BACKGROUND OF THE INVENTION

Field of the Invention

5 The invention relates to an implantable medical device such as an anastomosis device and a deployment system for implanting the device. In a preferred embodiment, the device can be used for forming a sutureless connection between a bypass graft and a blood vessel.

Brief Description of the Related Art

10 Vascular anastomosis is a procedure by which two blood vessels within a patient are surgically joined together. Vascular anastomosis is performed during treatment of a variety of conditions including coronary artery disease, diseases of the great and peripheral vessels, organ transplantation, and trauma. In coronary artery disease (CAD) an occlusion or stenosis in a coronary artery interferes with blood flow to the heart muscle. Treatment of CAD involves the grafting of a
15 vessel in the form of a prosthesis or harvested artery or vein to reroute blood flow around the occlusion and restore adequate blood flow to the heart muscle. This treatment is known as coronary artery bypass grafting (CABG).

20 In the conventional CABG, a large incision is made in the chest and the sternum is sawed in half to allow access to the heart. In addition, a heart lung machine is used to circulate the patient's blood so that the heart can be stopped and the anastomosis can be performed. During this procedure, the aorta is clamped which can lead to trauma of the aortic tissue and/or dislodge plaque emboli, both of which increase the likelihood of neurological complications. In order to minimize the trauma to the patient induced by conventional CABG, less invasive

techniques have been developed in which the surgery is performed through small incisions in the patients chest with the aid of visualizing scopes. Less invasive CABG can be performed on a beating or stopped heart and thus may avoid the need for cardiopulmonary bypass.

5 In both conventional and less invasive CABG procedures, the surgeon has to suture one end of the graft vessel to the coronary artery and the other end of the graft vessel to a blood supplying vein or artery. The suturing process is a time consuming and difficult procedure requiring a high level of surgical skill. In order to perform the suturing of the graft to the coronary artery and the blood supplying
10 artery the surgeon must have relatively unobstructed access to the anastomosis site within the patient. In the less invasive surgical approaches, some of the major coronary arteries including the ascending aorta cannot be easily reached by the surgeon because of their location. This makes suturing either difficult or impossible for some coronary artery sites. In addition, some target vessels, such
15 as heavily calcified coronary vessels, vessels having very small diameter, and previously bypassed vessels may make the suturing process difficult or impossible.

Accordingly, it would be desirable to provide a sutureless vascular anastomosis device which easily connects a graft to a target vessel. It would also be desirable to provide a sutureless anastomosis device which is formed of one
20 piece and is secured to the target vessel in a single step.

SUMMARY OF THE INVENTION

A superelastic or pseudoelastic one piece anastomosis device according to the present invention connects a graft vessel to a target vessel. The anastomosis device deforms from an insertion configuration to a tissue holding configuration
25 due to the superelastic or pseudoelastic properties of the material.

5

10

20

BRIEF DESCRIPTION OF THE DRAWINGS

The invention will now be described in greater detail with reference to the preferred embodiments illustrated in the accompanying drawings, in which like elements bear like reference numerals, and wherein:

5 FIG. 1 is a perspective view of a first embodiment of an anastomosis device in a constrained configuration prior to use;

 FIG. 2 is a perspective view of the anastomosis device of FIG. 1 in a deployed configuration;

 FIG. 3 is a side cross sectional view of the anastomosis device of FIG. 1
10 with a graft vessel everted around the device and the device constrained by a tube prior to deployment;

 FIG. 4 is a side cross sectional view of the system of FIG. 3 being inserted into a target vessel:

 FIG. 5 is a side cross sectional view of the system of FIG. 3 after release
15 of an inner flange;

 FIG. 6 is a side cross sectional view of the system of FIG. 3 with the inner flange imbedded in an inner wall of the target vessel wall;

 FIG. 7 is a side cross sectional view of the system of FIG. 3 after release of the outer flange showing the deployment tube being removed;

20 FIG. 8 is a perspective view of an alternative embodiment of an anastomosis device in a constrained configuration prior to use;

 FIG. 9 is a perspective view of the anastomosis device of FIG. 8 in a deployed configuration;

25 FIG. 10 is side cross sectional view of the anastomosis device of FIG. 8 after deployment shown connecting a graft and a target vessel;

 FIG. 11 is a perspective view of an alternative embodiment of an anastomosis device in a constrained configuration prior to use;

FIG. 12 is a perspective view of the anastomosis device of FIG. 11 in a deployed configuration;

FIG. 13 is a side cross sectional view of the anastomosis device of FIG. 11 after deployment shown connecting a graft vessel to a target vessel;

5 FIG. 14 is a perspective view of an alternative embodiment of an anastomosis device in a constrained configuration prior to use;

FIG. 15 is a perspective view of the anastomosis device of FIG. 14 in a deployed configuration;

10 FIG. 16 is a perspective view of an alternative embodiment of an expandable body anastomosis device in a constrained configuration prior to use; and

FIG. 17 is a perspective view of the anastomosis device of FIG. 16 in a deployed configuration.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

15 The present invention relates to a superelastic or pseudoelastic anastomosis device and method for connecting a graft vessel to a target vessel without the use of conventional sutures. The quick and easy deployment of the anastomosis system according to the present invention greatly increases the speed with which anastomosis can be performed over the known sutured anastomosis methods. The
20 anastomosis devices according to the present invention are particularly designed for use in connecting graft vessels to target vessel in a variety of anastomosis procedures, including coronary artery bypass grafting. In such procedures, a large vessel anastomotic device is used to connect a graft vessel to large diameter target vessels such as the aorta or it's major side branches and a small vessel anastomotic
25 device is used for connecting a graft vessel to a target vessel having a small diameter such as a coronary artery.

Suturing a graft vessel to a target vessel with conventional procedures is difficult and time consuming, particularly in minimally invasive procedures where space may be limited and in procedures in which it may be desired to perform an anastomosis without stoppage of blood flow through the target vessel. The superelastic or pseudoelastic anastomosis device and method of the present invention allow anastomosis to be performed efficiently and effectively in tight spaces. The anastomosis may also be performed with or without stoppage of blood flow in a target vessel and with or without the use of cardiopulmonary bypass.

FIG. 1 illustrates an anastomosis device 10 according to a first embodiment of the present invention in a constrained insertion configuration in which the anastomosis device would be inserted into a target blood vessel. FIG. 2 illustrates the anastomosis device 10 of FIG. 1 in an expanded deployed configuration which holds a graft vessel to a target vessel. The superelastic or the pseudoelastic anastomosis device in FIG. 1 includes a substantially cylindrical body 14, a plurality of legs 16 extending from a first side of the body, and a plurality of hooks 18 extended from a second side of the body. In the insertion configuration illustrated in FIG. 1, the body 14, the legs 16, and the hooks 18, are substantially aligned in a constrained substantially cylindrical shape. The anastomosis device 10 may be held in the constrained substantially cylindrical shape by a deployment tool, such as a substantially cylindrical deployment tube. When the deployment tool is removed from the device 10, the device returns to a preset expanded shape illustrated in FIG. 2 due to the superelastic or pseudoelastic properties of the material.

The anastomosis device 10 is made of a pseudoelastic or superelastic alloy, such as Nitinol or other pseudoelastic or superelastic material. The superelastic or pseudoelastic device 10 will self deform through superelastic or pseudoelastic

behavior from the constrained insertion configuration illustrated in FIG. 1 to the expanded configuration illustrated in FIG. 2 when the constraining device or deployment tool is removed. The anastomosis device 10 formed of the superelastic or pseudoelastic material is formed in the final shape illustrated in

5 FIG. 2 and is then isothermally deformed by constraining in a tube or other deployment tool in the substantially cylindrical shape illustrated in FIG. 1. The need for temperature control is avoided since the anastomosis device 10 reforms the deployed shape of FIG. 2 spontaneously when removed from the constraining tube. This allows the accurate placement of the anastomosis device 10

10 spontaneous and nearly instantaneously upon deployment of the device. The need for a mechanical deployment device to mechanically deform the anastomosis device from the insertion configuration to the deployed configuration is also avoided.

The anastomosis devices of the present invention may be made of any

15 known superelastic or pseudoelastic material. U.S. Patent No. 5,597,378 provides a discussion of superelastic and pseudoelastic materials and is incorporated herein by reference in its entirety.

The deployed anastomosis device 10 as shown in FIG. 2 includes an inner flange formed by outwardly extruding ends 22 of the J-shaped hooks 18. The

20 deployed device 10 also includes an outer flange formed by the legs 16 extending outward from the body 14.

In use, a graft vessel 30, shown in FIG. 3, is threaded through a center of the anastomosis device 10. An end 34 of the graft vessel 30 is everted around the hooks 18 and the hook ends 22 penetrate into or through the everted end 34 of the

25 graft vessel retaining the graft vessel in place on the anastomosis device 10.

As illustrated in FIG. 3, the anastomosis device 10 with the everted graft vessel 30 is positioned within a deployment tube 38 for delivery of the anastomosis

device and graft vessel to an opening 40 in a target vessel 32. In the radially constrained insertion configuration, the leading edge or hook end of the anastomosis device may be substantially cylindrical or slightly conical for ease of insertion.

5 One embodiment of a method for deploying the anastomosis device 10 of the present invention will be described with reference to FIGS. 3-6. As shown in FIG. 3, the graft vessel 30 is prepared by everting an end 34 of the graft vessel around the hooks 18 of the anastomosis device 10. The hook ends 22 penetrate the graft vessel tissue to maintain the everted configuration of the graft vessel. The
10 hooks 18 and the legs 16 of the anastomosis device 10 are radially constrained by inserting the anastomosis device 10 and everted end of the graft vessel 30 into a deployment tool 38 in the shape of a tube. When positioned inside the deployment tool 38, the anastomosis device 10 is in a generally cylindrical configuration for insertion into the target vessel 32.

15 As shown in FIG. 4, the deployment tool 38 is used to insert the anastomosis device 10 and the graft vessel 30 into the target vessel 32 until the hook ends 22 have passed through the opening 40 and are positioned within an interior of the blood vessel. As shown in FIGS. 3 and 4, a retainer tube 36 is positioned around the graft vessel 30 and inside the deployment tool 38 for holding
20 and extruding the anastomosis device 10. A distal end 48 of the retainer tube 46 is positioned adjacent to a proximal end of the anastomosis device 10. The distal end 48 of the retainer tube 46 may be attached to or abut the anastomosis device 10 to hold the anastomosis device in place inside the deployment tube 38 during the insertion step of FIG. 4.

25 As shown in FIG. 5, the anastomosis device 10 is held in place by the retainer tube 46 while the deployment tube 38 is withdrawn or retracted to release the radial constraining force from the hooks 18. Upon removal of the deployment

tube 38 from the hooks 18, the hook ends 22 and hook base portion 24 spontaneously spring outward due to the superelasticity or pseudoelasticity of the material.

As shown in FIG. 6, after the release of the hooks 18 the anastomosis device 10 is withdrawn by the deployment tool 38 against the interior wall of the target vessel 32 causing the hook ends 22 to be compressed against or penetrate into the tissue of the interior wall of the target vessel. The deployment tube 38 is then completely withdrawn as shown in FIG. 7 allowing the legs 16 to spontaneously spring outward to trap the wall of the target vessel 32 between the hooks 18 which form an inner flange and the legs 16 which form an outer flange for the deployed anastomosis device 10.

FIGS. 8-10 illustrate an alternative embodiment of an anastomosis device 50 having a central body portion 54. A first set of legs 56 extend from one end of the body 54 and a second set of pointed legs 58 extend from the second side of the body. In a constrained configuration illustrated in FIG. 8, the anastomosis device 50 is substantially tubular for insertion into a target vessel. In an expanded deployed configuration, illustrated in FIGS. 9 and 10, the anastomosis device 50 is substantially C-shaped in cross section with the legs 56 forming an outer flange and the pointed legs 58 forming an inner flange of the anastomosis device 50.

The embodiment shown in FIGS. 8-10 may be deployed in a manner similar to that of the anastomosis device described above with respect to FIGS. 1-7. As shown in FIG. 10, the graft vessel 62 is everted around the anastomosis device 50. The anastomosis device 50 and graft vessel 62 are then inserted into an opening in the target vessel 64 in a constrained configuration. A constraining device such as the deployment tool 38 is then removed from the anastomosis device 50 and graft vessel 62 allowing the legs 56 and 58 to spontaneously spring outward by the superelastic or pseudoelastic properties of the material to form

inner and outer flanges which trap the tissue of the target vessel 64 between the inner and outer flanges.

According to one preferred embodiment of the anastomosis device 50 the pointed legs 58 each include a pointed tissue penetrating end 66 and a rectangular stop member 68 for limiting the tissue penetration of the penetrating end. As shown in FIG. 10, the tissue penetrating end 66 of the pointed legs 58 penetrates into or through the graft vessel 62 to ensure the graft vessel is retained on the anastomosis device 50 during and after deployment.

In the deployed configuration illustrated in FIG. 10, the intima of the graft vessel 62 abuts an intima of the target vessel 64. Thus, the expansion of the inner flange of the anastomosis device 50 forms a vein gasket to seal the graft and target vessels together.

FIGS. 11-13 illustrate an alternative embodiment of the superelastic or pseudoelastic anastomosis device 80 in a radially constrained configuration illustrated in FIG. 11 and in an expanded tissue retaining configuration illustrated in FIGS. 12 and 13. The anastomosis device 80 includes a device body 84 formed of a plurality of substantially parallel spring elements 86 interconnecting to end members 88. Extending from the end members 88 are a plurality of prongs 90 which in the expanded tissue supporting configuration illustrated in FIG. 13, form inner and outer flanges to trap the tissue of the target vessel 96. As in the previous embodiments, a graft vessel 94 is inserted through a center of the anastomosis device body 84 and is everted around the prongs 90 of at least one end of the device body. The prongs 90 penetrate into or through the graft vessel tissue to retain the graft vessel on the anastomosis device.

The anastomosis device 80 with the graft vessel 94 everted around the anastomosis device is inserted in a radially constrained configuration illustrated in FIG. 11 into an opening in the target vessel 96. When the radially constraining

member such as a retainer tube is removed from the anastomosis device 80, the anastomosis device spontaneously self deforms and returns to the configuration of FIG. 12 due to the superelastic or pseudoelastic properties of the material.

As shown in FIG. 13, a first set of the prongs 90 forms a flange at the inner wall of the target vessel. The spring elements 86 allow the distance between the end members 88 to adjust somewhat to target vessels 96 having walls of different thicknesses. The spring elements 86 may also apply a compression force to the wall of the target vessel 96 once the anastomosis device 80 has been deployed to provide improved sealing.

In an alternate embodiment of the anastomosis device 80 of FIGS. 11-13, the graft vessel 94 may be attached to the anastomosis device without everting. This may be done by providing axial prongs, hooks, or barbs on the inner rail member 88 and hooking an end of the graft vessel on the hooks, prongs, or barbs without everting.

An alternative embodiment of an anastomosis device 100 includes an anastomosis device body 104, legs 106, and hooks 108, as in the embodiment of FIGS. 1 and 2. The embodiment of FIGS. 14 and 15 differs from the embodiment of FIGS. 1 and 2 in that the legs 106 are folded outward and downward adjacent the body 104 in the radially constrained insertion configuration illustrated in FIG. 14. The legs 106 will spontaneously spring out to the flange forming configuration of FIG. 15 when the radially constraining member such as a retainer tube is removed for deployment of the anastomosis device 100.

FIGS. 16 and 17 illustrate an alternative embodiment of an anastomosis device 200 including a device body 204, legs 206 and pointed legs 208. The body 204 is formed of axially extending members 210 interconnected by struts 212 which allow the body to expand radially. Positioned between the body 204 and the pointed legs 208 are hinges 214. FIG. 16 illustrates the anastomosis device 200

in a radially constrained insertion configuration with a graft vessel 220 extending through an interior of the device body 204 and everted over the pointed legs 208. The pointed legs 208 penetrate and hold the everted end of the graft vessel 220 on the device 200.

5 For insertion, the anastomosis device 200 of FIG. 16 is radially constrained in a deployment tube (not shown). As the deployment tube is withdrawn from the device 200, the pointed legs 208 fold outward to form an inner flange, the device body 204 expands radially, and the legs 206 fold outward to form an outer flange. The radially expanding body 204 helps to stretch and support an opening in the target vessel.

10 Each of the anastomosis devices according to the present invention are preferably single piece devices which are formed in a substantially tubular shape. The anastomosis devices may be formed by laser cutting or punching from a tube or sheet of superelastic or pseudoelastic material. Alternatively, the devices may be formed from superelastic or pseudoelastic wire. The devices may be provided in varying sizes to join vessels of different sizes. The legs, hooks, prongs, and other device elements which have been discussed above with regard to the various embodiments may be used in varying numbers and arrangements depending on the particular application.

15 20 The invention has been described as an anastomosis device which is constrained for insertion in a radially constrained configuration with a deployment tool such as tube. However, the deployment tube may take other non-tubular shapes.

25 Although the invention has been primarily discussed with respect to coronary artery bypass surgery, the anastomosis devices of the present invention may be used in other types of anastomosis procedures. For example, the anastomosis device may be used in femoral-femoral bypass, vascular shunts,

subclavian-carotid bypass, organ transplants, and the like. The devices according to the present invention may be used with venous grafts such as a harvested saphenous vein graft, arterial graft, such as a dissected mammal artery, or a synthetic prosthesis, as required.

5 Finally, the anastomosis devices according to the present invention have been illustrated as substantially cylindrical members. However, the devices can also be shaped into ovals, football shapes, or other shapes. Oval shapes can be particularly useful for accommodating small target vessels.

10 While the invention has been described in detail with reference to the preferred embodiments thereof, it will be apparent to one skilled in the art that various changes and modifications can be made and equivalents employed, without departing from the present invention.

6. The device of Claim 4, wherein the tissue penetrating elements extend radially outwardly from the device body for holding an everted end of the graft vessel.

1 7. The device of Claim 1, wherein the device body uses the
2 superelastic or pseudoelastic properties of the material to self deform from the
3 insertion configuration to the tissue holding configuration.

1 8. A tube deployed anastomosis system for connecting a graft vessel to
2 a target vessel comprising:
3 a deployment tube; and
4 an anastomosis device formed of a superelastic or pseudoelastic
5 material, the device having an insertion configuration and a tissue holding
6 configuration in which the device has an inner flange and an outer flange, wherein
7 the inner and outer flanges are radially constrained in the deployment tube in the
8 insertion configuration for insertion into the target vessel and when released from
9 the deployment tube, the device self deforms to the tissue holding configuration.

1 9. The device of Claim 8, wherein a portion of the device body
2 between the inner flange and the outer flange is expandable from a first diameter
3 insertion configuration to a second diameter tissue holding configuration.

1 10. The device of Claim 8, wherein the superelastic or pseudoelastic
2 material is a nickel titanium alloy.

1 11. The device of Claim 8, further comprising a plurality of tissue
2 penetrating elements for penetrating and holding a graft vessel in place on the
3 device body.

1 12. The device of Claim 11, wherein the tissue penetrating elements
2 extend radially outwardly from the device body for holding an everted end of the
3 graft vessel.

1 13. The device of Claim 8, wherein the device body uses the
2 superelastic or pseudoelastic properties of the material to self deform from the
3 insertion configuration to the tissue holding configuration.

1 14. A method of deploying an anastomosis system for connecting a graft
2 vessel to a target vessel, the method comprising:
3 connecting a graft vessel to a one piece device formed of a
4 superelastic or pseudoelastic material;
5 poking a portion of the one piece device through the graft vessel;
6 and
7 deploying the one piece device by self deformation to a tissue
8 holding configuration in which the device has an inner flange and an outer flange
9 and traps the target vessel tissue between the inner flange and the outer flange.

1 15. The method of Claim 14, wherein the one piece device is deployed
2 by removing a radially constraining deployment tool from the device.

1 16. The method of Claim 15, wherein the deployment tool is a
2 deployment tube which receives the tubular device, and wherein the deployment
3 tube is inserted partially into the target vessel wall and then withdrawn to deploy
4 the one piece device from the deployment tube.

1 17. The method of Claim 14, wherein the one-piece device is deployed
2 by employing the superelastic or pseudoelastic property of a superelastic or
3 pseudoelastic material from which the one piece device is formed.

1 18. The method of Claim 14, wherein the graft vessel is everted around
2 the one piece device.

1 19. The method of Claim 14, wherein the deployed one piece device
2 abuts an intima of the graft vessel against an intima of the target vessel.

ABSTRACT OF THE DISCLOSURE

A one piece anastomosis device is disclosed which is formed of a superelastic or pseudoelastic material which self deforms or self deploys from an insertion configuration to a tissue holding configuration. The device in a deployed state preferably includes an inner tissue penetrating flange which penetrate and
5 retains an everted graft vessel and an outer flange. The self deploying anastomosis device does not rely on a temperature transformation to achieve deployment.

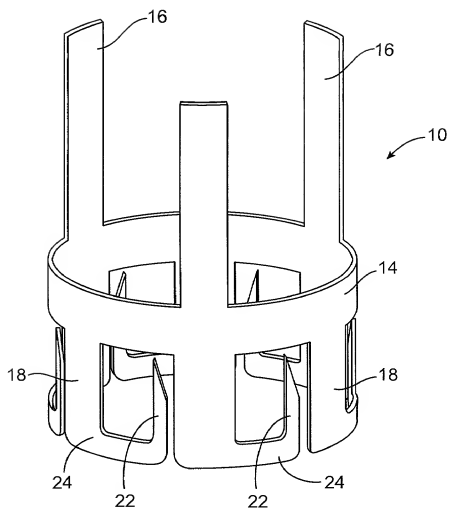


FIG. 1

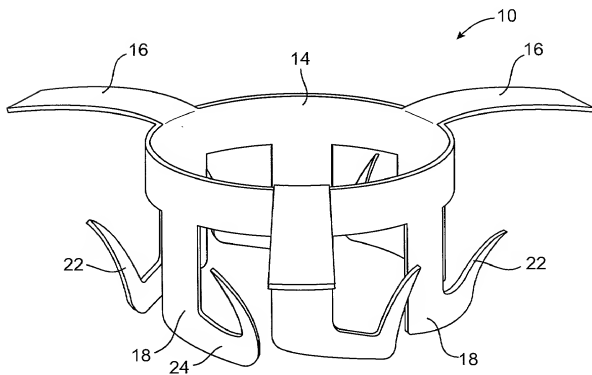


FIG. 2

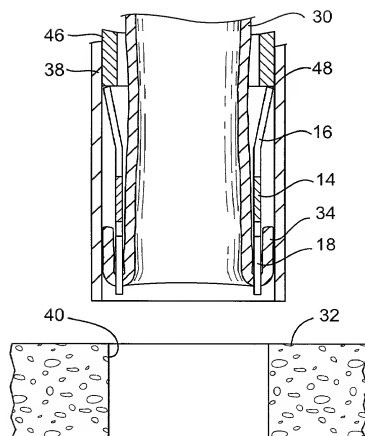


FIG. 3

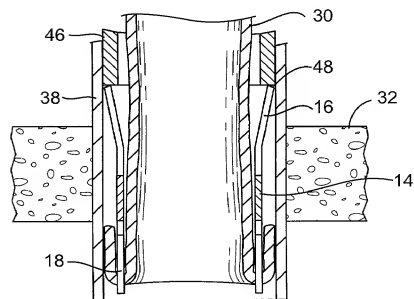
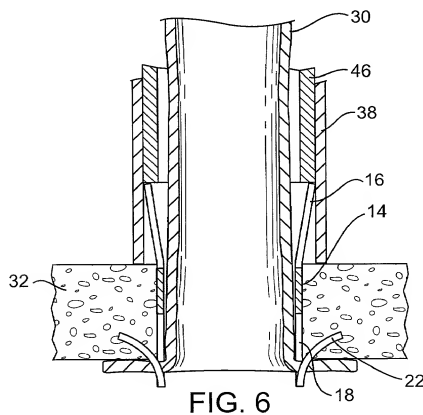
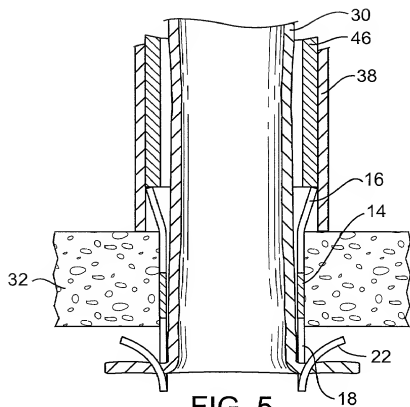


FIG. 4



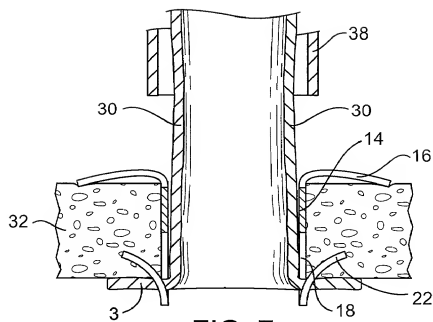


FIG. 7

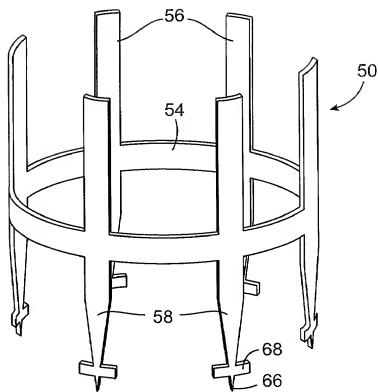


FIG. 8

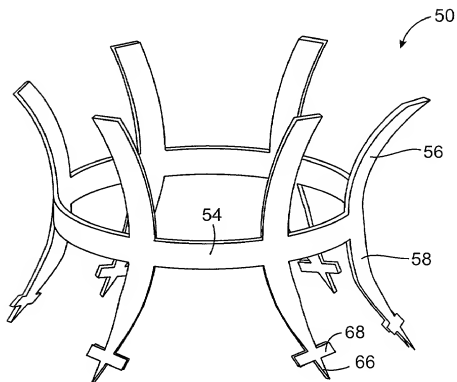


FIG. 9

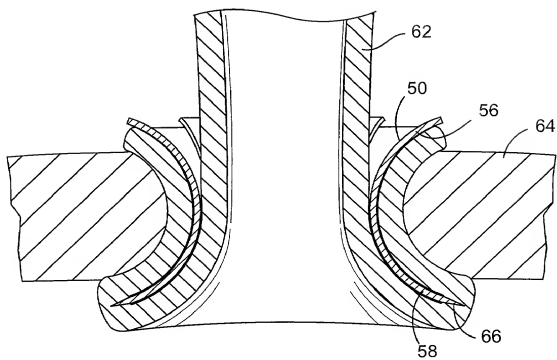


FIG. 10

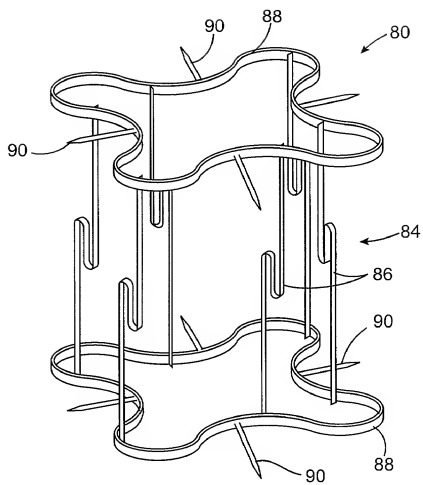


FIG. 11

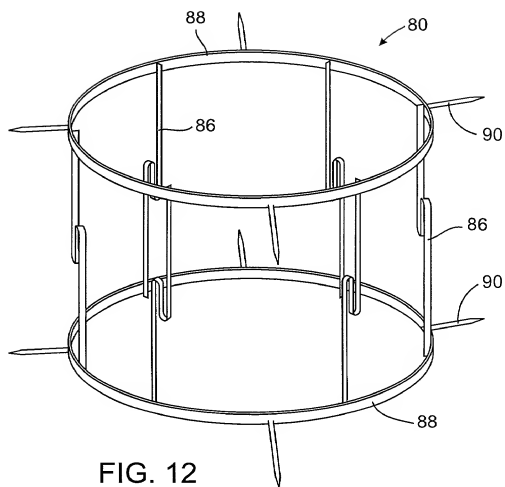


FIG. 12

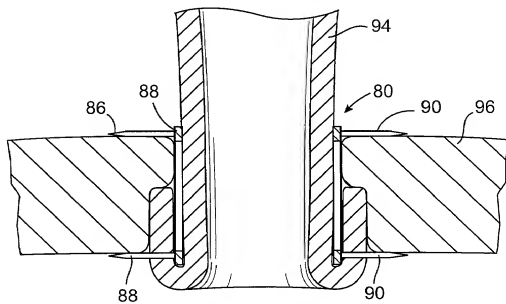


FIG. 13

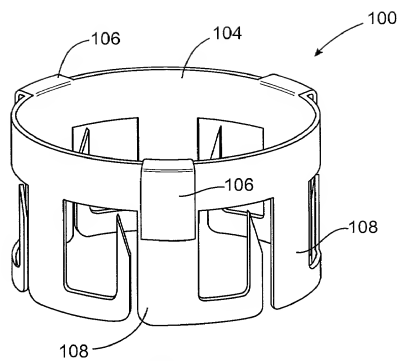


FIG. 14

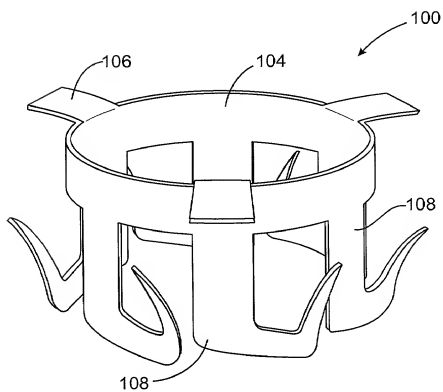


FIG. 15

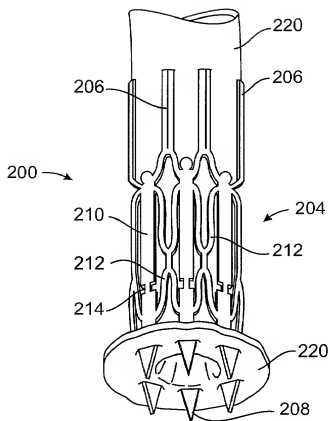


FIG. 16

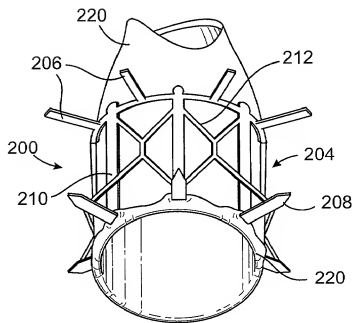


FIG. 17

**COMBINED DECLARATION AND POWER OF ATTORNEY
FOR UTILITY PATENT APPLICATION**

Attorney's Docket No.

032405-018

As a below-named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name;

I BELIEVE I AM THE ORIGINAL, FIRST AND SOLE INVENTOR (if only one name is listed below) OR AN ORIGINAL, FIRST AND JOINT INVENTOR (if more than one name is listed below) OF THE SUBJECT MATTER WHICH IS CLAIMED AND FOR WHICH A PATENT IS SOUGHT ON THE INVENTION ENTITLED:

IMPLANTABLE SUPERELASTIC ANASTOMOSIS DEVICE

the specification of which

(check one)



is attached hereto;



was filed on _____ as

Application No. _____

and was amended on _____;
(if applicable)

I HAVE REVIEWED AND UNDERSTAND THE CONTENTS OF THE ABOVE-IDENTIFIED SPECIFICATION, INCLUDING THE CLAIMS, AS AMENDED BY ANY AMENDMENT REFERRED TO ABOVE;

I ACKNOWLEDGE THE DUTY TO DISCLOSE TO THE OFFICE ALL INFORMATION KNOWN TO ME TO BE MATERIAL TO PATENTABILITY AS DEFINED IN TITLE 37, CODE OF FEDERAL REGULATIONS, Sec. 1.56 (as amended effective March 16, 1992);

I do not know and do not believe the said invention was ever known or used in the United States of America before my or our invention thereof, or patented or described in any printed publication in any country before my or our invention thereof or more than one year prior to said application; that said invention was not in public use or on sale in the United States of America more than one year prior to said application; that said invention has not been patented or made the subject of an inventor's certificate issued before the date of said application in any country foreign to the United States of America on any application filed by me or my legal representatives or assigns more than twelve months prior to said application;

I hereby claim foreign priority benefits under Title 35, United States Code Sec. 119 and/or Sec. 365 of any foreign application(s) for patent or inventor's certificate as indicated below and have also identified below any foreign application for patent or inventor's certificate on this invention having a filing date before that of the application(s) on which priority is claimed:

COMBINED DECLARATION AND POWER OF ATTORNEY

Attorney's Docket No.

032405-018

COUNTRY/INTERNATIONAL

APPLICATION NUMBER

DATE OF FILING
(day, month, year)PRIORITY
CLAIMED

YES_ NO_

YES_ NO_

I hereby appoint the following attorneys and agent(s) to prosecute said application and to transact all business in the Patent and Trademark Office connected therewith and to file, prosecute and to transact all business in connection with international applications directed to said invention:

| | | | | | |
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| Robert S. Swecker | 19,885 | Eric H. Weisblatt | 30,505 | Charles F. Wieland III | 33,096 |
| Platon N. Mandros | 22,124 | James W. Peterson | 26,057 | Bruce T. Wiedler | 33,815 |
| Benton S. Duffett, Jr. | 22,030 | Teresa Stanek Rea | 30,427 | Todd R. Walters | 34,040 |
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

FULL NAME OF SOLE OR FIRST INVENTOR

SIGNATURE

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